This Page Is Inserted by IFW Operations and is not a part of the Official Record

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

IMAGES ARE BEST AVAILABLE COPY.

As rescanning documents will not correct images, please do not report the images to the Image Problem Mailbox.

(43) Application published 6 Nov 1985

- (21) Application No 8505279
- (22) Date of filing 1 Mar 1985
- (30) Priority data
 - (31) 8405521
- (32) 2 Mar 1984
- (33) GB
- (71) Applicant
 A B M Chemicals Limited (United Kingdom),
 Unity Mills, Poleacre Lane, Woodley, Stockport, Cheshire
- (72) inventors
 Dr Roy Hatton,
 Dr Peter Lees,
 Brian Davis
- (74) Agent and/or Address for Service Stevens Hewlett & Perkins, 5 Quality Court, Chancery Lane, London WC2A 1HZ

- (51) INT CL⁴ C08G 65/32 C11D 1/722 3/075
- (52) Domestic classification
 C3R 27K8E J
 C5D 6A5B 6A5D2 6A5E 6B11A 6B12H 6B12N2 6B1 6B2
 6C6 6C8
 U1S 1379 1427 C3R C5D
- (56) Documents cited None
- (58) Field of search C2C C3R

(54) End-blocked nonionic surfactants

(57) Alkylbenzyl poly(oxyalkylene) derivatives of the formula

wherein R_1 is a straight or branched chain 1–4C alkyl group; R_2 is a straight or branched chain 4–20C alkyl group; R_3 and R_4 are each, independently, H or CH_3 ; n and m are each integers wherein n + m does not exceed 20 have good nonionic surface activity, biodegradability and are stable to caustic powders. The compounds are useful in alkaline cleaning compositions, such as dishwater products, and also in rinse-aid formulations for use in automatic dishwashers.

10

15

20

30

35

40

60

45

60

End-blocked nonionic surfactants

5 The present invention relates to end-blocked nonionic surfactants. More particularly, it relates to new alkylbenzyl poly(oxyalkylene) derivatives useful as nonionic surfactants.

In the past, a number of classes of compounds have been used, in detergents and janitorial compositions, as non-ionic surface active agents. Examples of these include amine ethoxylate propoxylates, benzylblocked alkylphenol alkoxylates and ethylene oxide-propylene oxide block copolymers. These materials suffer the major disadvantage that they lack biodegradability and are unstable on solid caustic soda. Furthermore, such materials as are commercially available are ill-defined or have incompletely characterisable chemical structure.

In an attempt to overcome the problem of lack of biodegradability of prior nonionic surfactants, alcohol ethoxylate propoxylates, having molecular weights less than 1000, were used. Unfortunately, these materials have poor dispersibility in water at temperatures above their cloud points. In addition, they are unstable in the presence of solid caustic powders, e.g. caustic soda, as in the case of the earlier nonionics.

Futher attempts to overcome the drawbacks of the prior art materials involved "blocking" the terminal hydroxyl group in fatty alcohol ethoxylates with alkyl chlorides. The procedures used, however, were difficult and they made use of toxic and expensive materials. Similarly blocked non-ionics using benzyl chloride as alkylating agent were introduced although the commercial product was only about 70% blocked, indicating the poor efficiency of the blocking procedure.

We have now discovered a range of compounds having good nonionic surface activity which are biodegradable and stable to caustic powders. Furthermore, we have discovered that these compounds can be prepared by a process which achieves greater than 90% blocking.

The present invention provides new compounds which are represented by the following general formula I.

$$R_1$$
-0(CH₂CHO)_n (CH₂CHO)—CH₂ R₂

wherein R₁ is a straight or branched chain alkyl group having from 1 to 4 carbon atoms; R₂ is a straight or branched chain alkyl group having from 4 to 20 carbon atoms; R₃ and R₄ are each, independently, H or -CH₃, and n and m are each integers wherein the sum n + m is not greater than 20.

As we have stated above, the group R₁ is a 1 to 4 C alkyl, such as methyl, ethyl, propyl, isopropyl, n-butyl, 35 isobutyl, sec-butyl and tert-butyl. The alkyl group R₂ attached to the phenyl ring of the blocking group is a straight or branched chain group containing from 4 to 20 carbon atoms, preferably 8 to 12 carbon atoms. Preferred examples of suitable R₂ groups include straight and branched chain octyl, decyl and dodecyl groups. A particularly preferred R₂ group is n-dodecyl.

The sum n+m is not greater than 20 and will usually lie within the range of from 5 to 15, more particularly 40 within the range 8 to 14. A preferred value of n+m is 10.

Some preferred compounds of the present invention are as follows:

Me
$$O(CH_2CH_2O)_{10} CH_2 - C_8H_{17}$$

25

30

35

40

45

50

55

60

2

20

25

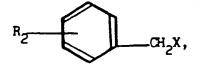
30

35

5

$$_{10}$$
 n-Bu 0 (CH₂CH₂0)₆CH₂ $c_{12}^{H}_{25}$

The present invention also provides a process for preparing a compound of the formula I above which process comprises reacting a lowr alcohol of the formula R₁OH, wherein R₁ is as defined above, with an alkylene oxide selected from ethylene oxide and propylene oxide, and then reacting the ethoxylated and/or propoxylated material with an alkylbenzyl halide having the formula



wherein X is a halogen atom, preferably a chlorine atom. In a preferred embodiment, the ethoxylated and/or propoxylated material is reacted with a mixture of alkylbenzyl chlorides, such as a mixture of C_{θ} and C_{10} alkylbenzyl chlorides to give a mixture of reaction products.

If it is desired to form a compound of formula I wherein R₃ is H and R₄ is -CH₃, then the lower alcohol is first 40 reacted with ethylene oxide and the ethoxylated product is then reacted with propylene oxide. If the desired compound of formula I has R_3 = CH_3 and R_4 =H, then the alcohol is first propoxylated and the alkyl propoxylate is then ethoxylated. The amount of reactants used in this step, of course, depend upon the desired values of n and m in the final compound. Usually, the amounts of alkylene oxide used will be sufficient such that the sum of n + m is within the range of from 5 to 15. The alkoxylated lower alcohol is then 45 condensed with the alkylbenzyl chloride. The condensation reaction will usually be conducted in the presence of an alkali metal hydroxide, typically NaOH. The condensation reaction is preferably phase transfer catalysed with, for instance, a catalytic amount of a tetraalkyl ammonium halide. By the use of this process, it is possible to achieve a blocking efficiency of greater than 90%. Furthermore, it should be noticed that, in the above-described process, end-blocking of the molecule and the introduction of the hydrophobe 50 into the molecule are carried out in one step unlike prior art processes. Thus, by the process of the invention, it is possible to make end blocked non ionics at low cost.

The compounds of the invention have relatively low molecular weight and meet E.E.C. legislation on bi degradability (Directive 82/242/EEC). They are stable on solid caustic powders and have excellent def aming properties, go d wetting and detergency. The compounds fth invention are, theref re, 55 particularly well suited to use in strongly alkaline cleaning c mp sitlons in combinati n with s lid caustic p wders, such as p wdered caustic soda. In particular, the compounds f the present inventin are especially useful as I w-foaming ranti-foaming ingredients in alkaline powder dishwashing products. They may also be used advantage usly in rinse-aid formulati ns for automatic dishwashing and may c ntrol or pr vent f am generated by r stabilis d by food soils.

Example 1

60

1 mold of n-butanol was ethoxylated with 10 m les fethylene oxide by a conventional meth d. The eth xylated product (150 g) was then charged together with 50% caustic s da solution (80 g) and, as catalyst, benzyltriethylammonium chloride (2.5 g). The temperature of this mixture was raised to 90-95°C at which postabling a mixture of p-octyl and p-decylbenzyl chloride (45 a) was run in over a period of 3 h lurs. The

65

30

40

60

20

reaction mixture was then heated to 110°C and maintained at that t mp rature for 1 hour. After cooling, the reaction mixture was washed with water to remove salt, unreacted polyglycol and catalyst. The product comprised a mixture of:-

Example 2

The procedure of Example 1 was repeated, except that methanol, instead of n-butanol, was used. The product comprised a mixture of

25
$$Me0(CH_2CH_20)_{10}CH_2 - C_{10}H_{21}$$
 25

Example 3
The procedure of Example 1 was repeated except that dodecyl benzyl chloride, instead of p-octyl and p-decyl

30 benzyl chloride, was used.
The product comprised

Some of the properties of the products obtained in Examples 1, (A), and 2, (B), were measured and compared with those obtained for a prior art unblocked non-ionic surfactant, nonyl phenol 9.5 mole ethoxylate (NP 95). These are set out below.

			Α	В	NP95	
45	1.	Cloud point, at concentration of 1% in water, in °C	34	47	52	45
	2.	Foam (by Ross Miles), at a concentration of 0.1% in water and temperature 5°C below				
		cloud point, in mm.	41	61	100	
50	3.	Foam at cloud point, at concentration of 0.1% in				50
		water, in mm.	30	28	66	
	4.	Wetting (by Draves) at a concentrati n of 0.1% in				
55		water and temperature 5°C below cloud point, in seconds	11	21	4	55
	5.	Stability on caustic	good	good	very poor.	

These results ship with the compositions A and B, which are in accordance with thip resent invention, show a big foam reduction compared to the primary compound and suffer only small loss in wetting proporties.

The compounds of this invention are advantage usly used as low-foaming or antifoaming ingredients in alkalin powder dishwashing products.

10

25

30

35

40

60

Example 4

:0

40

An alkaline powder dishwashing product with good cleaning p wer and controlled foaming characteristics is prepared by blending the following:

5	Sodium metasilicate	30% 38%	by weight by weight	
	Soda ash *Trisodium nitrilo triacetic acid powder	10%	by weight	
	Sodium tripolyphosphate	20%	by weight	
	Suitable compound of present invention	2%	by weight	

*e.g. Nervanaid NTA Conc. Powder, a trade name of A.B.M. Chemicals Ltd.

wherein the suitable compound of the present invention is one or a mixture of more than one of the following:

a) A compound according to formula 1 wherein (n+m)=10 and $R_3=R_4=H \text{ and } R_1=C_2H_5 \text{ and } R_2=C_{12}H_{25}$ b) a compound according to formula 1 wherein (n+m)=6 and $R_3=R_4=H \text{ and } R_1=C_4H_9 \text{ (n-butyl) and } R_2=C_{12}H_{25}.$ 20

c) a compound according to formula 1 wherein

 $(n \div m) = 14 \text{ and}$

 $R_3 = R_4 = H \text{ and } R_1 = C_4H_9 \text{ (n-butyl)}$ and $R_2 = C_{12}H_{25}$.

The compounds of the invention are advantageously used in rinse-aid formulations for automatic dishwashing. Spotless quick-drying is achieved without causing subsequent foaming problems in those machines where rinse water is re-cycled into the wash-water. Furthermore formulations

containing compounds of the present invention may control or prevent foam generated or stabilised by food soils.

Example 5

Rinse aid formulations for automatic dishwashing according to the following recipes give spotless quick-drying:

Suitable compound of present invention 25%
Dipropylene glycol 10%
N-lauroyl sarcosinate 1%
Water 64%

Where a suitable compound of the present invention includes a mixture of compounds according to formula 1, wherein, for example, (n + m) = 6 and 14 and $R_3 = R_4 = H$ and $R_1 = C_4H_9$ (n-butyl) and $R_2 = C_{12}H_{25}$. Such mixture comprises 60% of such compound wherein (n + m) = 14 and 40% of such compound wherein (n + m) = 6, and when the formulation of Example 5 is made with this mixture of non-ionic surface active agents a formulation (Formulation 1) with properties superior to those of current commercial formulations is obtained, as measured in the following tests:

50		Actives % (Drying at 110°C)	Protein* Foam (mm)	Haze Temp. (°C)	Dispersibility in water at 80℃	Glass rinse appearanc at 80°C	50
	Formulation 1	36	24	61	Excellent	Excellent	
55	Commercial Product A	30	184	51	Fair	Very good	55
	Commercial Product B	50	50	60	G od	Fair	-
	Commercial Pr duct C	32	90	47	G od	Fair ·	

Protein foam tests measur the height if foam priduced in a dynamic foaming test with 100 ppm of rinse aid in water of 150 ppm hardness in the presence of 0.25% milk protein and 0.36% alkaline builders at a temperature if 60°C. The test involes pumping the test solution at a pressure of 5 psi through a 1.8 mm in rifice with a fall of 750 mm ont the surface if the test solution with recirculation for 5 min.

15

20

35

55

25

30

40

CLAIMS

1. A compound which is represented by the following general formula:-

$$R_{10}(CH_{2}CHO) = CH_{2}CHO) = CH_{2}CHO$$

- 10 wherein R₁ is a straight or branched chain 1-4C alkyl group; R₂ is a straight or branched chain 4-20C alkyl group; R₃ and R₄ are each, independently, H or −CH₃; and n and m are each integers wherein the sum n + m is not greater than 20.
 - 2. A compound as claimed in claim 1, wherein R₂ is a straight or branched chain alkyl group having from 8 to 12 carbon atoms.
- 15 3. A compound as claimed in claim 2, wherein R₂ is a straight or branched chain octyl, decyl or dodecyl group.
 - 4. A compound as claimed in any one of claims 1 to 3, wherein the sum of n + m is in the range of from 5 to 15.
- 5. A compound as claimed in any one of claims 1 to 4, wherein R_1 is n-butyl; R_2 is a substantially linear 20 dodecyl; R_3 and R_4 are both H and the sum n + m is in the range of from 5 to 15.
 - 6. A compound as claimed in any one of claims 1 to 4, wherein R_1 is ethyl; R_2 is a substantially linear dodecyl; R_3 and R_4 are both H and the sum n + m is in the range of from 5 to 15.
 - 7. A method of preparing a compound having the following general formula:-

$$R_1O(CH_2CHO)_{\overline{n}}$$
 $(CH_2CHO)_{\overline{m}}$ CH_2 R_2 R_2

wherein R_1 is a straight or branched chain 1-4C alkyl group; R_2 is a straight or branched chain 4-20C alkyl group; R_3 and R_4 are each, independently, H or $-CH_3$; and n and m are each integers wherein the sum n + m is not greater than 20,

which method comprises reacting an alochol of the formula R₁OH, where R₁ is as defined above, with an alkylene oxide selected from ethylene oxide, propylene oxide and mixtures thereof and then reacting the resulting alkoxylated material with an alkylbenzyl halide having the formula

wherein R₂ is as defined above and X is a halogen atom.

8. A method of preparing a compound having the following general formula:-

R₁0(CH₂CH₂)
$$\frac{CH_3}{n}$$
 CH₂CH₂CH₂ $\frac{CH_3}{m}$ $\frac{CH_2}{n}$ $\frac{R_2}{n}$ 50

wherein R₁ is a straight or branched chain 1-4C alkyl group; R₂ is a straight or branched chain 4-20C alkyl group; and n and m ar each integ rs wher in the sum n + m is not greater than 20, which method comprises reacting an alochol of the formula R₁OH, where R₁ is as defined above, with ethylene oxide, then reacting the resulting ethoxylated product with propylene oxide and then reacting the resulting propoxylated material with an alkylbenzyl halide of the formula:—

60

É

15

5

10

25

where R_2 is as defined abov and X is a halogen atom.

9. A method of preparing a compound having the following general formula:-

wherein R₁ is a straight or branched chain 1-4C alkyl group; R₂ is a straight or branched chain 4-20C alkyl group; and n and m are each integers wherein the sum n + m is not greater than 20, which method comprises reacting an alcohol of the formula R₁OH, where R₁ is as defined above, with propylene oxide, reacting the resulting propoxylated material with ethylene oxide and then reacting the resulting ethoxylated product with an alkylbenzyl halide of the formula

R₂ CH Y

20 wherein R₂ is as defined above and X is a halogen atom.

10. A cleaning composition comprising a mixture of caustic soda and at least one compound as claimed in claim 1.

11. A cleaning composition substantially as hereinbefore described in Example 4.

25 12. A rinse aid formulation for use in an automatic dishwater substantially as hereinbefore described in Example 5.

Printed in the UK for HMSO, D8818935, 9/85, 7102.
Published by The Patent Office, 25 Southampton Buildings, London, WC2A 1AY, from which copies may be obtained.

.